

Compounds of the formula I in which G represents -CH₂NR¹⁶-, -CONR¹⁶, CH₂N(R¹⁶)-T- or -CH₂N(R¹⁶)COT- wherein R¹⁶ is not hydrogen, may be prepared from the appropriate compound of the formula I wherein R¹⁶ is hydrogen by introducing the appropriate R¹⁶ by acylation, alkylation etc. For example, by using similar methods to those disclosed in the specific examples.

IN THE CLAIMS:

Please cancel claims 1, 3, 10-12, and 14-17 from the present application without disclaimer or prejudice.

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Please amend claims 7, 8, 9, and 13 as follows:

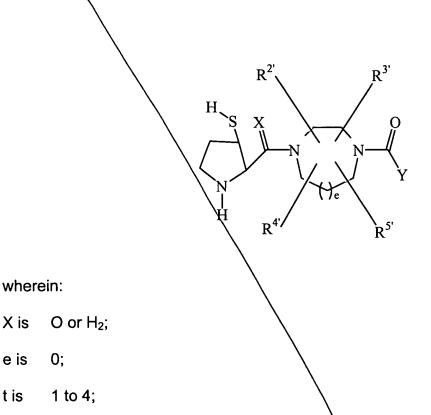
Claim 7:

X is

e is

t is

A compound of the formula B: 7.



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R^{2'}, R^{3'}, R^{4'}, and R^{5'} are independently selected from: H; C₁₋₈alkyl, alkenyl, alkynyl, aryl, heterocycle, -CO-NR^{6'}R^{7'} or -CO-OR^{6'}, unsubstituted or substituted with one or more of:

- 1) ary or heterocycle, unsubstituted or substituted with:
 - a. $\backslash C_{1-4}$ alkyl,
 - b. $(\dot{C}H_2)tOR^{6'}$,
 - c. $(CH_{\lambda})tNR^{6'}R^{7'}$,
 - d. halogen,
- 2) C₃₋₆cycloalkyl,
- 3) OR^{6} ,
- 4) SR^{6′}, S(O)R^{6′}, SO₂R^{6′},
- 5) $-NR^{6'}R^{7'}$,
- 6) $-NR^{6'}-CO-R^{7'}$,
- 7) $-NR^{6'}-CO-NR^{7'}R^{8'}$,
- 8) $-O-CO-NR^{6'}R^{7'}$,
- 9) -O-CO-OR⁶,
- 10) -O-NR⁶'R⁷',
- 11) -SO₂NR^{6′}R^{7′},
- 12) $-NR^{6'}-SO_2-R^{7'}$,
- 13) -CO-R⁶, or
- 14) -CO-OR⁶;

and any two of R^{2'}, R^{3'}, R^{4'}, and R^{5'} are optionally attached to the same carbon atom;

Y is aryl, heterocycle, unsubstituted or substituted with one or more of:

1) C₁₋₄alkyl, unsubstituted or substituted with:

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& DUNNER, L. L. P.
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- α . C_{1-4} alkoxy,
 - $b NR^{6'}R^{7'}$
- c. \setminus C₃₋₆cycloalkyl,
- d. aryl or heterocycle,
- е. HQ,
- 2) aryl or heterocycle,
- 3) halogen,
- 4) OR⁶',
- 5) $NR^{6'}R^{7'}$,
- 6) CN
- 7) NO_2 , or
- 8) CF₃;

 $R^{6'}$, $R^{7'}$ and $R^{8'}$ are independently selected from: H; $C_{1\text{--}4}$ alkyl, $C_{3\text{--}6}$ cycloalkyl, heterocycle, aryl, aroyl, heteroaroyl, arylsulfonyl, heteroarylsulfonyl, unsubstituted or substituted with:

- a) C₁₋₄alkoxy,
- b) aryl or heterocycle,
- c) halogen,
- d) HO,
- e) -CO-R^{9'},
- f) -SO₂R^{9'}, wherein

R^{6'} and R^{7'} may be joined in a ring, and

R^{7'} and R^{8'} may be joined in a ring;

R^{9'} is C₁₋₄alkyl or aralkyl;

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a pharmaceutically acceptable salt thereof.

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Claim 8:

8. The compound (2<u>S</u>)-2-(2-methoxyethyl)-1-((cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-4-naphthoyl-piperazine or a pharmaceutically acceptable salt thereof.

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Claim 9:

9. A pharmaceutical composition which comprises a compound according to any one of claims 7 or 8 and a pharmaceutically acceptable carrier.



Claim 13:

13. A process for preparing compounds of the Formula B as defined in claim 7 which comprises deprotecting a compound of Formula VI:

wherein X⁸ represents the right hand side of the Formula B as defined in claim 7, Pr¹ is H or an amino protecting group, Pr² is H or a thio protecting group and any functional groups in X⁸ are optionally protected with the proviso that there is at least one protecting group and optionally, if desired, converting the product thus obtained into a pharmaceutically acceptable salt thereof.

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& DUNNER, L. L. P.
1300 I STREET, N. W.
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Please add the following claims:



- 18. A method of treating a disease or medical condition mediated through farnesylation of CAAX-containing proteins which comprises administering to a warm-blooded animal an effective amount of a compound according to claim 7 or 8, wherein said disease or medication condition is a carcinoma of the bladder, breast, colon, kidney, liver, lung, ovary, pancreas, stomach, cervix, thyroid or skin.
- 19. A method of treating a disease or medical condition mediated through farnesylation of CAAX-containing proteins which comprises administering to a warm-blooded animal an effective amount of a compound according to claim 7 or 8, wherein said disease or medical condition is a hematopoietic tumor of lymphoid lineage selected from acute lymphocytic leukaemia, B-cell lymphoma and Burketts lymphoma.
- 20. A method of treating a disease or medical condition mediated through farnesylation of CAAX-containing proteins which comprises administering to a warm-blooded animal an effective amount of a compound according to claim 7 or 8, wherein said disease or medical condition is a hematopoietic tumor of myeloid lineage selected from acute or chronic myelogenous leukemias and promyelocytic leukaemia.
- 21. A method of treating a disease or medical condition mediated through farnesylation of CAAX-containing proteins which comprises administering to a warm-blooded animal an effective amount of a compound according to claim 7 or 8, wherein said disease or medical condition is a tumor of mesenchymal origin selected from fibrosarcoma and rhabdomyosarcoma.
- 22. A method of treating a disease or medical condition mediated through farnesylation of CAAX-containing proteins which comprises administering to a warm-

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& DUNNER, L.L.P.
1300 I STREET, N.
WASHINGTON, DC 20005
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